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***** Welcome to STN International *****

NEWS	1		Web Page for STN Seminar Schedule - N. America
NEWS	2	NOV 21	CAS patent coverage to include exemplified prophetic substances identified in English-, French-, German-, and Japanese-language basic patents from 2004-present
NEWS	3	NOV 26	MARPAT enhanced with FSORT command
NEWS	4	NOV 26	CHEMSAFE now available on STN Easy
NEWS	5	NOV 26	Two new SET commands increase convenience of STN searching
NEWS	6	DEC 01	ChemPort single article sales feature unavailable
NEWS	7	DEC 12	GBFULL now offers single source for full-text coverage of complete UK patent families
NEWS	8	DEC 17	Fifty-one pharmaceutical ingredients added to PS
NEWS	9	JAN 06	The retention policy for unread STNmail messages will change in 2009 for STN-Columbus and STN-Tokyo
NEWS	10	JAN 07	WPIDS, WPINDEX, and WPIX enhanced Japanese Patent Classification Data
NEWS	11	FEB 02	Simultaneous left and right truncation (SLART) added for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
NEWS	12	FEB 02	GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS	13	FEB 06	Patent sequence location (PSL) data added to USGENE
NEWS	14	FEB 10	COMPENDEX reloaded and enhanced
NEWS	15	FEB 11	WTEXTILES reloaded and enhanced
NEWS	16	FEB 19	New patent-examiner citations in 300,000 CA/CAPLUS patent records provide insights into related prior art
NEWS	17	FEB 19	Increase the precision of your patent queries -- use terms from the IPC Thesaurus, Version 2009.01
NEWS	18	FEB 23	Several formats for image display and print options discontinued in USPATFULL and USPAT2
NEWS	19	FEB 23	MEDLINE now offers more precise author group fields and 2009 MeSH terms
NEWS	20	FEB 23	TOXCENTER updates mirror those of MEDLINE - more precise author group fields and 2009 MeSH terms
NEWS	21	FEB 23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS	22	FEB 25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS	23	MAR 06	INPADOCDB and INPAFAMDB enhanced with new display formats
NEWS	24	MAR 11	EPFULL backfile enhanced with additional full-text applications and grants
NEWS	25	MAR 11	ESBIOBASE reloaded and enhanced
NEWS	26	MAR 20	CAS databases on STN enhanced with new super role for nanomaterial substances
NEWS	27	MAR 23	CA/CAPLUS enhanced with more than 250,000 patent equivalents from China

NEWS 28 MAR 30 IMSPATENTS reloaded and enhanced

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 12:06:59 ON 30 MAR 2009

=> file medline embase biosis caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.22	0.22

FILE 'MEDLINE' ENTERED AT 12:07:20 ON 30 MAR 2009

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=> s Porchet H?/AU
L1 261 PORCHET H?/AU

=> s l1 and prostate
L2 7 L1 AND PROSTATE

=> dup rem l2
PROCESSING COMPLETED FOR L2
L3 4 DUP REM L2 (3 DUPLICATES REMOVED)

=> s Heimgarter F?/AU
L4 0 HEIMGARTER F?/AU

=> s Heimgartner F?/AU
L5 32 HEIMGARTNER F?/AU

=> s l5 and prostate
L6 2 L5 AND PROSTATE

=> dup rem l6
PROCESSING COMPLETED FOR L6

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L7          2 DUP REM L6 (0 DUPLICATES REMOVED)

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L8          29 CURDY C?/AU

=> s l8 and prostate
L9          3 L8 AND PROSTATE

=> dup rem l9
PROCESSING COMPLETED FOR L9
L10         3 DUP REM L9 (0 DUPLICATES REMOVED)

=> s Ducrey B?/AU
L11         16 DUCREY B?/AU

=> dup rem l11
PROCESSING COMPLETED FOR L11
L12         10 DUP REM L11 (6 DUPLICATES REMOVED)

=> s l12 and prostate
L13         3 L12 AND PROSTATE

=> dis his

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(FILE 'HOME' ENTERED AT 12:06:59 ON 30 MAR 2009)

FILE 'MEDLINE, EMBASE, BIOSIS, CAPLUS' ENTERED AT 12:07:20 ON 30 MAR 2009

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L1          261 S PORCHET H?/AU
L2          7 S L1 AND PROSTATE
L3          4 DUP REM L2 (3 DUPLICATES REMOVED)
L4          0 S HEIMGARTER F?/AU
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L8          29 S CURDY C?/AU
L9          3 S L8 AND PROSTATE
L10         3 DUP REM L9 (0 DUPLICATES REMOVED)
L11         16 S DUCREY B?/AU
L12         10 DUP REM L11 (6 DUPLICATES REMOVED)
L13         3 S L12 AND PROSTATE

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L3 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:1487237 CAPLUS

DOCUMENT NUMBER: 150:24287

TITLE: Slow release pharmaceutical composition made of microparticles comprising PLGA type copolymer and peptide salt as active substance

INVENTOR(S): Ducrey, Bertrand; Garrouste, Patrick; Curdy, Catherine; Bardet, Marie-Anne; Porchet, Herve ; Lundstrom, Eija; Heimgartner, Frederic

PATENT ASSIGNEE(S): Debio Recherche Pharmaceutique S.A., Switz.

SOURCE: PCT Int. Appl., 19pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2008149320 A2 20081211 WO 2008-IB52241 20080606
 WO 2008149320 A3 20090205

W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
 RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: EP 2007-109767 A 20070606
 WO 2007-IB54372 A 20071027

AB Pharmaceutical composition made of microparticles for the slow release of an active substance at least during a period covering the sixth month after injection of said composition, said composition comprising a group of microparticles

made of a copolymer of the PLGA type which incorporate an active substance in the form of a water insol. peptide salt; said copolymer furthermore comprising at least 75% of lactic acid and an inherent viscosity between 0.1 and 0.9 dL/g, as measured in chloroform at 25° and at a polymer concentration of 0.5 g/dL; said microparticles furthermore having a size distribution defined as follows : D(v,0.1) is between 10 and 30 µm, D(v,0.5) is between 30 and 70 µm, D(v,0.9) is between 50 and 110 µm.

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS ON SIN

ACCESSION NUMBER: 2005:612107 CAPLUS
 DOCUMENT NUMBER: 143:120559
 TITLE: Sustained release compositions of gonadotropin hormone releasing hormone and estrogen
 INVENTOR(S): Porchet, Herve; Lundstrom, Eija; Curdy, Catherine; Bardet, Marie-Anne
 PATENT ASSIGNEE(S): Debiopharm S.A., Switz.
 SOURCE: PCT Int. Appl., 30 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005063276	A1	20050714	WO 2004-IB4276	20041223
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

PRIORITY APPLN. INFO.: WO 2003-IB6159 A 20031223

AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns.

of the invention can be employed for an improved androgen deprivation therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flushes are minimized through the maintenance of a minimally adequate estrogen level. Combinations of polyestradiol phosphate and triptorelin were prepared for extemporaneous reconstitution.

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:965090 CAPLUS

DOCUMENT NUMBER: 141:389284

TITLE: Methods and compositions using gonadotropin hormone releasing hormone

INVENTOR(S): Porchet, Herve; Helmgartner, Frederic;

Curdy, Catherine; Ducrey, Bertrand

PATENT ASSIGNEE(S): Debiopharm S.A., Switz.

SOURCE: PCI Int. Appl., 33 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004096259	A1	20041111	WO 2004-IB1334	20040430
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2523830	A1	20041111	CA 2004-2523830	20040430
EP 1617859	A1	20060125	EP 2004-730606	20040430
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
BR 2004009950	A	20060425	BR 2004-9950	20040430
CN 1780634	A	20060531	CN 2004-80011409	20040430
JP 2006525306	T	20061109	JP 2006-506555	20040430
MX 2005011299	A	20060124	MX 2005-11299	20051020
US 20070042040	A1	20070222	US 2006-554292	20060922
PRIORITY APPLN. INFO.:			WO 2003-IB1680	A 20030430
			WO 2004-IB1334	W 20040430

AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns.

of the invention can be employed for an improved androgen deprivation therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flashes are minimized through the maintenance of a minimally adequate estrogen level.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 4 MEDLINE on STN

ACCESSION NUMBER: 2003355146 MEDLINE

DUPLICATE 1

DOCUMENT NUMBER: PubMed ID: 12887472
 TITLE: Comparative efficacy of triptorelin pamoate and leuprolide acetate in men with advanced prostate cancer.
 AUTHOR: Heyns C F; Simonin M-P; Grossgurin P; Schall R; Porchet H C
 CORPORATE SOURCE: Department of Urology, University of Stellenbosch, Tygerberg Hospital, Western Cape, South Africa. (For the South African Triptorelin Study Group). cfh2@sun.ac.za
 SOURCE: BJU international, (2003 Aug) Vol. 92, No. 3, pp. 226-31. Journal code: 100886721. ISSN: 1464-4096.
 PUB. COUNTRY: England; United Kingdom
 DOCUMENT TYPE: (CLINICAL TRIAL)
 (COMPARATIVE STUDY)
 Journal; Article; (JOURNAL ARTICLE)
 (MULTICENTER STUDY)
 (RANDOMIZED CONTROLLED TRIAL)
 (RESEARCH SUPPORT, NON-U.S. GOV'T)
 LANGUAGE: English
 FILE SEGMENT: Priority Journals
 ENTRY MONTH: 200309
 ENTRY DATE: Entered STN: 31 Jul 2003
 Last Updated on STN: 5 Sep 2003
 Entered Medline: 4 Sep 2003
 AB OBJECTIVE: To compare the efficacy of monthly administrations of the luteinizing hormone-releasing hormone agonists triptorelin pamoate and leuprolide acetate to induce and maintain castrate levels of serum testosterone in men with advanced prostate cancer. PATIENTS AND METHODS: Men with advanced prostate cancer were randomly assigned to receive triptorelin 3.75 mg or leuprolide 7.5 mg. The agent was injected intramuscularly every 28 days for nine injections. Primary endpoints were the percentages of men whose serum testosterone concentrations declined to and were maintained at or below castrate levels (≤ 1.735 nmol/L or ≤ 500 ng/L) during 9 months (253 days) of treatment. Secondary endpoints were luteinizing hormone levels, bone pain, prostate specific antigen levels, quality of life, testosterone pharmacodynamics, survival, and safety variables. RESULTS: In all, 284 men received either triptorelin (140) or leuprolide (144). The percentage of men with castrate levels of serum testosterone was lower at 29 days for triptorelin than for leuprolide (91.2% vs 99.3%; point estimate - 8.0, 95% confidence interval - 16.9% to - 1.4%), but equivalent at 57 days (97.7% vs 97.1%). The mean (98.8% vs 97.3%) and cumulative (96.2% vs 91.2%) castration maintenance rates between 29 and 253 days were equivalent between the treatment groups. Secondary endpoints were equivalent between treatment groups except for the 9-month survival rate, which was significantly higher for triptorelin than for leuprolide (97.0% vs 90.5%; $P = 0.033$). Both treatments were well tolerated. CONCLUSION: Triptorelin reduced testosterone concentrations less rapidly, but maintained castration as effectively as leuprolide. There was no evidence that the slower onset of castration caused deleterious effects.

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	ENTRY	SESSION
FULL ESTIMATED COST	36.82	37.04
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-2.46	-2.46

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FILE CONTAINS CURRENT INFORMATION.

LAST RELOADED: Mar 27, 2009 (20090327/UP).

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FILE 'MEDLINE, EMBASE, BIOSIS, CAPLUS' ENTERED AT 12:07:20 ON 30 MAR 2009

L1 261 S PORCHET H7/AU
L2 7 S L1 AND PROSTATE
L3 4 DUP REM L2 (3 DUPLICATES REMOVED)
L4 0 S HEIMGARTNER F7/AU
L5 32 S HEIMGARTNER F7/AU
L6 2 S L5 AND PROSTATE
L7 2 DUP REM L6 (0 DUPLICATES REMOVED)
L8 29 S CURDY C7/AU
L9 3 S L8 AND PROSTATE
L10 3 DUP REM L9 (0 DUPLICATES REMOVED)
L11 16 S DUCREY B7/AU
L12 10 DUP REM L11 (6 DUPLICATES REMOVED)
L13 3 S L12 AND PROSTATE

FILE 'STNGUIDE' ENTERED AT 12:11:28 ON 30 MAR 2009

=> dis ibib abs l6 1-2

YOU HAVE REQUESTED DATA FROM FILE 'CAPLUS' - CONTINUE? (Y)/N:y

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:1487237 CAPLUS

DOCUMENT NUMBER: 150:24287

TITLE: Slow release pharmaceutical composition made of microparticles comprising PLGA type copolymer and peptide salt as active substance

INVENTOR(S): Ducrey, Bertrand; Garrouste, Patrick; Curdy, Catherine; Bardet, Marie-Anne; Porchet, Herve; Lundstrom, Eija; Heimgartner, Frederic

PATENT ASSIGNEE(S): Debio Recherche Pharmaceutique S.A., Switz.

SOURCE: PCT Int. Appl., 19pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2008149320	A2	20081211	WO 2008-IB52241	20080606
WO 2008149320	A3	20090205		
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RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK,			

TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
 TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,
 AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: EP 2007-109767 A 20070606
 WO 2007-IB54372 A 20071027

AB Pharmaceutical composition made of microparticles for the slow release of an active substance at least during a period covering the sixth month after injection of said composition, said composition comprising a group of microparticles made of a copolymer of the PLGA type which incorporate an active substance in the form of a water insol. peptide salt; said copolymer furthermore comprising at least 75% of lactic acid and an inherent viscosity between 0.1 and 0.9 dL/g, as measured in chloroform at 25° and at a polymer concentration of 0.5 g/dL; said microparticles furthermore having a size distribution defined as follows : D(v,0.1) is between 10 and 30 µm, D(v,0.5) is between 30 and 70 µm, D(v,0.9) is between 50 and 110 µm.

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:965090 CAPLUS

DOCUMENT NUMBER: 141:389284

TITLE: Methods and compositions using gonadotropin hormone releasing hormone

INVENTOR(S): Porchet, Herve; Heimgartner, Frederic;
 Curdy, Catherine; Ducrey, Bertrand

PATENT ASSIGNEE(S): Debiopharm S.A., Switz.

SOURCE: PCT Int. Appl., 33 pp.
 CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004096259	A1	20041111	WO 2004-IB1334	20040430
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CA 2523830	A1	20041111	CA 2004-2523830	20040430
EP 1617859	A1	20060125	EP 2004-730606	20040430
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BR 2004009950	A	20060425	BR 2004-9950	20040430
CN 1780634	A	20060531	CN 2004-80011409	20040430
JP 2006525306	T	20061109	JP 2006-506555	20040430
MX 2005011299	A	20060124	MX 2005-11299	20051020
US 20070042040	A1	20070222	US 2006-554292	20060922
PRIORITY APPLN. INFO.:			WO 2003-IB1680 A 20030430	
			WO 2004-IB1334 W 20040430	
AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns. of the invention can be employed for an improved androgen deprivation				

therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flashes are minimized through the maintenance of a minimally adequate estrogen level.
 REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> dis his

(FILE 'HOME' ENTERED AT 12:06:59 ON 30 MAR 2009)

FILE 'MEDLINE, EMBASE, BIOSIS, CAPLUS' ENTERED AT 12:07:20 ON 30 MAR 2009

L1 261 S PORCHET H7/AU
 L2 7 S L1 AND PROSTATE
 L3 4 DUP REM L2 (3 DUPLICATES REMOVED)
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 L5 32 S HEIMGARTNER F7/AU
 L6 2 S L5 AND PROSTATE
 L7 2 DUP REM L6 (0 DUPLICATES REMOVED)
 L8 29 S CURDY C7/AU
 L9 3 S L8 AND PROSTATE
 L10 3 DUP REM L9 (0 DUPLICATES REMOVED)
 L11 16 S DUCREY B7/AU
 L12 10 DUP REM L11 (6 DUPLICATES REMOVED)
 L13 3 S L12 AND PROSTATE

FILE 'STNGUIDE' ENTERED AT 12:11:28 ON 30 MAR 2009

FILE 'CAPLUS' ENTERED AT 12:12:35 ON 30 MAR 2009

FILE 'STNGUIDE' ENTERED AT 12:12:35 ON 30 MAR 2009

=> dis ibib abs l10 1-3

YOU HAVE REQUESTED DATA FROM FILE 'CAPLUS' - CONTINUE? (Y)/N:y

L10 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2008:1487237 CAPLUS
 DOCUMENT NUMBER: 150:24287
 TITLE: Slow release pharmaceutical composition made of microparticles comprising PLGA type copolymer and peptide salt as active substance
 INVENTOR(S): Ducrey, Bertrand; Garrouste, Patrick; Curdy, Catherine; Bardet, Marie-Anne; Porchet, Herve; Lundstrom, Eija; Heimgartner, Frederic
 PATENT ASSIGNEE(S): Debio Recherche Pharmaceutique S.A., Switz.
 SOURCE: PCT Int. Appl., 19pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2008149320	A2	20081211	WO 2008-IB52241	20080606
WO 2008149320	A3	20090205		
W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE,				

KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
 RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: EP 2007-109767 A 20070606
 WO 2007-IB54372 A 20071027

AB Pharmaceutical composition made of microparticles for the slow release of an active substance at least during a period covering the sixth month after injection of said composition, said composition comprising a group of microparticles

made of a copolymer of the PLGA type which incorporate an active substance in the form of a water insol. peptide salt; said copolymer furthermore comprising at least 75% of lactic acid and an inherent viscosity between 0.1 and 0.9 dL/g, as measured in chloroform at 25° and at a polymer concentration of 0.5 g/dL; said microparticles furthermore having a size distribution defined as follows : D(v,0.1) is between 10 and 30 µm, D(v,0.5) is between 30 and 70 µm, D(v,0.9) is between 50 and 110 µm.

L10 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:612107 CAPLUS

DOCUMENT NUMBER: 143:120559

TITLE: Sustained release compositions of gonadotropin hormone releasing hormone and estrogen

INVENTOR(S): Porchet, Herve; Lundstrom, Eija; Curdy, Catherine; Bardet, Marie-Anne

PATENT ASSIGNEE(S): Debiopharm S.A., Switz.

SOURCE: PCT Int. Appl., 30 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005063276	A1	20050714	WO 2004-IB4276	20041223
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: WO 2003-IB6159 A 20031223

AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns.

of the invention can be employed for an improved androgen deprivation therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flushes are minimized through the maintenance of a minimally adequate estrogen level. Combinations of polyestradiol phosphate and triptorelin were prepared for

extemporaneous reconstitution.
REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2004:965090 CAPLUS
DOCUMENT NUMBER: 141:389284
TITLE: Methods and compositions using gonadotropin hormone
releasing hormone
INVENTOR(S): Porchet, Herve; Heimgartner, Frederic; Curdy,
Catherine; Ducrey, Bertrand
PATENT ASSIGNEE(S): Debiopharm S.A., Switz.
SOURCE: PCT Int. Appl., 33 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004096259	A1	20041111	WO 2004-IB1334	20040430
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2523830	A1	20041111	CA 2004-2523830	20040430
EP 1617859	A1	20060125	EP 2004-730606	20040430
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
BR 2004009950	A	20060425	BR 2004-9950	20040430
CN 1780634	A	20060531	CN 2004-80011409	20040430
JP 2006525306	T	20061109	JP 2006-506555	20040430
MX 2005011299	A	20060124	MX 2005-11299	20051020
US 20070042040	A1	20070222	US 2006-554292	20060922
PRIORITY APPLN. INFO.:			WO 2003-IB1680	A 20030430
			WO 2004-IB1334	W 20040430

AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns. of the invention can be employed for an improved androgen deprivation therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flashes are minimized through the maintenance of a minimally adequate estrogen level.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> dis his

(FILE 'HOME' ENTERED AT 12:06:59 ON 30 MAR 2009)

FILE 'MEDLINE, EMBASE, BIOSIS, CAPLUS' ENTERED AT 12:07:20 ON 30 MAR 2009
L1 261 S PORCHET H7/AU

L2 7 S L1 AND PROSTATE
 L3 4 DUP REM L2 (3 DUPLICATES REMOVED)
 L4 0 S HEIMGARTER F?/AU
 L5 32 S HEIMGARTNER F?/AU
 L6 2 S L5 AND PROSTATE
 L7 2 DUP REM L6 (0 DUPLICATES REMOVED)
 L8 29 S CURDY C?/AU
 L9 3 S L8 AND PROSTATE
 L10 3 DUP REM L9 (0 DUPLICATES REMOVED)
 L11 16 S DUCREY B?/AU
 L12 10 DUP REM L11 (6 DUPLICATES REMOVED)
 L13 3 S L12 AND PROSTATE

FILE 'STNGUIDE' ENTERED AT 12:11:28 ON 30 MAR 2009

FILE 'CAPLUS' ENTERED AT 12:12:35 ON 30 MAR 2009

FILE 'STNGUIDE' ENTERED AT 12:12:35 ON 30 MAR 2009

FILE 'CAPLUS' ENTERED AT 12:13:17 ON 30 MAR 2009

FILE 'STNGUIDE' ENTERED AT 12:13:17 ON 30 MAR 2009

=> dis ibib abs l13 1-3

YOU HAVE REQUESTED DATA FROM FILE 'EMBASE, CAPLUS' - CONTINUE? (Y)/N:y

L13 ANSWER 1 OF 3 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN
 ACCESSION NUMBER: 1994361901 EMBASE
 TITLE: [Current research on medicinal plants].
 RECHERCHE ACTUELLE DANS LE DOMAINE DES PLANTES MEDICINALES.
 AUTHOR: Ducrey, B. (correspondence)
 CORPORATE SOURCE: Insti. de Pharmacognosie/Phytochimie, Universite de Lausanne, BEP, 1015 Lausanne-Dorigny, Switzerland.
 SOURCE: Schweizer Apothekerzeitung, (1994) Vol. 132, No. 24, pp. 646-649.
 ISSN: 0036-7508 CODEN: SAPOEF
 COUNTRY: Switzerland
 DOCUMENT TYPE: Journal; Conference Article; (Conference paper)
 FILE SEGMENT: 016 Cancer
 030 Clinical and Experimental Pharmacology
 037 Drug Literature Index
 004 Microbiology: Bacteriology, Mycology, Parasitology and Virology
 LANGUAGE: French
 SUMMARY LANGUAGE: French
 ENTRY DATE: Entered STN: 29 Dec 1994
 Last Updated on STN: 29 Dec 1994

L13 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2008:1487237 CAPLUS
 DOCUMENT NUMBER: 150:24287
 TITLE: Slow release pharmaceutical composition made of microparticles comprising PLGA type copolymer and peptide salt as active substance
 INVENTOR(S): Ducrey, Bertrand; Garrouste, Patrick; Curdy, Catherine; Bardet, Marie-Anne; Porchet, Herve; Lundstrom, Eija; Heimgartner, Frederic
 PATENT ASSIGNEE(S): Debio Recherche Pharmaceutique S.A., Switz.

SOURCE: PCT Int. Appl., 19pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008149320	A2	20081211	WO 2008-IB52241	20080606
WO 2008149320	A3	20090205		
W:	AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
PRIORITY APPLN. INFO.:			EP 2007-109767 A 20070606	
			WO 2007-IB54372 A 20071027	

AB Pharmaceutical composition made of microparticles for the slow release of an active substance at least during a period covering the sixth month after injection of said composition, said composition comprising a group of microparticles

made of a copolymer of the PLGA type which incorporate an active substance in the form of a water insol. peptide salt; said copolymer furthermore comprising at least 75% of lactic acid and an inherent viscosity between 0.1 and 0.9 dL/g, as measured in chloroform at 25° and at a polymer concentration of 0.5 g/dL; said microparticles furthermore having a size distribution defined as follows : D(v,0.1) is between 10 and 30 µm, D(v,0.5) is between 30 and 70 µm, D(v,0.9) is between 50 and 110 µm.

L13 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2009 ACS ON SIN

ACCESSION NUMBER: 2004:965090 CAPLUS
 DOCUMENT NUMBER: 141:389284
 TITLE: Methods and compositions using gonadotropin hormone releasing hormone
 INVENTOR(S): Porchet, Herve; Heimgartner, Frederic; Curdy, Catherine; Ducrey, Bertrand
 PATENT ASSIGNEE(S): Debiopharm S.A., Switz.
 SOURCE: PCT Int. Appl., 33 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004096259	A1	20041111	WO 2004-IB1334	20040430
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,			

AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
 EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
 SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
 SN, TD, TG

CA 2523830	A1	20041111	CA 2004-2523830	20040430
EP 1617859	A1	20060125	EP 2004-730606	20040430
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
BR 2004009950	A	20060425	BR 2004-9950	20040430
CN 1780634	A	20060531	CN 2004-80011409	20040430
JP 2006525306	T	20061109	JP 2006-506555	20040430
MX 2005011299	A	20060124	MX 2005-11299	20051020
US 20070042040	A1	20070222	US 2006-554292	20060922

PRIORITY APPLN. INFO.:

WO 2003-IB1680	A	20030430
WO 2004-IB1334	W	20040430

AB The present invention relates to compns. comprising two sustained release formulations, the first being capable of releasing a gonadotropin releasing hormone composition and the second an estrogenic composition The compns.

of the invention can be employed for an improved androgen deprivation therapy of prostate cancer, in which therapy loss of bone mineral d. and the occurrence and severity of hot flashes are minimized through the maintenance of a minimally adequate estrogen level.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
0.07	65.91

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
0.00	-8.20

CA SUBSCRIBER PRICE

STN INTERNATIONAL LOGOFF AT 12:14:25 ON 30 MAR 2009